WHAT IS CLAIMED IS:

1	1.	A method of detecting a breast cancer-associated transcript in a cen		
2	from a patient, the method comprising contacting a biological sample from the patient with a			
3	polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence			
4	as shown in Tables 1-25.			
1	2.	The method of claim 1, wherein the biological sample comprises		
1	isolated nucleic acid			
2	Isolated fluciele act	uo.		
1	3.	The method of claim 2, wherein the nucleic acids are mRNA.		
1	4.	The method of claim 2, further comprising the step of amplifying		
2	nucleic acids before the step of contacting the biological sample with the polynucleotide.			
1	5.	The method of claim 1, wherein the polynucleotide comprises a		
2	sequence as shown	in Tables 1-25.		
1	6.	The method of claim 1, wherein the polynucleotide is immobilized on		
2	a solid surface.			
1	7.	The method of claim 1, wherein the patient is undergoing a therapeutic		
2	regimen to treat breast cancer.			
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1	8.	The method of claim 1, wherein the patient is suspected of having		
2	breast cancer.			
1	9.	An isolated nucleic acid molecule consisting of a polynucleotide		
2	sequence as shown			
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1	10.	The nucleic acid molecule of claim 9, which is labeled.		
1	11.	An expression vector comprising the nucleic acid of claim 9.		
1	12	A host cell comprising the expression vector of claim 11.		

1		13.	An isolated polypeptide which is encoded by a nucleic acid molecule
2	having polynu	cleotid	e sequence as shown in Tables 1-25.
.1		14.	An antibody that specifically binds a polypeptide of claim 13.
1		15.	The antibody of claim 14, further conjugated to an effector component
1		16.	The antibody of claim 15, wherein the effector component is a
2	fluorescent lab	el.	
1		17.	The antibody of claim 15, wherein the effector component is a
<u>_</u> 2	radioisotope o	r a cyto	otoxic chemical.
22		18.	The antibody of claim 15, which is an antibody fragment.
		19.	The antibody of claim 15, which is a humanized antibody
1 1 2 - 3		20.	A method of detecting a breast cancer cell in a biological sample from
2	_	nethod	comprising contacting the biological sample with an antibody of claim
1 3	14.		
1		21.	The method of claim 20, wherein the antibody is further conjugated to
2	an effector co	mpone	nt.
1		22.	The method of claim 21, wherein the effector component is a
2	fluorescent la	bel.	
1		23.	A method for identifying a compound that modulates a breast cancer-
2	associated po	lypepti	de, the method comprising the steps of:
3		(i) co	ntacting the compound with a breast cancer-associated polypeptide, the
4	polypeptide e	ncoded	by a polynucleotide that selectively hybridizes to a sequence at least
5	80% identical	to a se	equence as shown in Tables 1-25; and
6		(ii) de	etermining the functional effect of the compound upon the polypeptide.
. 1		24.	A drug screening assay comprising the steps of

(1)	administering a test compound to a mammal having breast cancer of a cen-			
isolated therefrom;				
(ii) comparing the level of gene expression of a polynucleotide that selectively			
hybridizes to a se	equence at least 80% identical to a sequence as shown in Tables 1-25 in a			
treated cell or mammal with the level of gene expression of the polynucleotide in a control				
cell or mammal,	wherein a test compound that modulates the level of expression of the			
polynucleotide is	a candidate for the treatment of breast cancer.			